

PCT

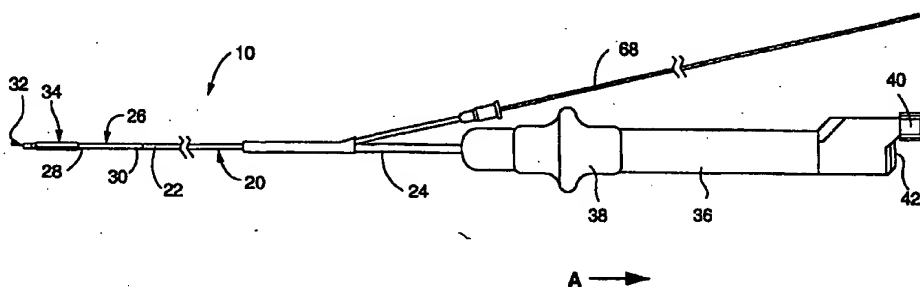
WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification <sup>6</sup> : <b>A61B 5/042, A61N 1/05, G01B 7/28, A61M 25/00, 25/01</b>		A1	(11) International Publication Number: <b>WO 99/48421</b> (43) International Publication Date: 30 September 1999 (30.09.99)
(21) International Application Number: PCT/US99/00567 (22) International Filing Date: 11 January 1999 (11.01.99) (30) Priority Data: 09/049,332 27 March 1998 (27.03.98) US (71) Applicant: CARDIAC PATHWAYS CORPORATION [US/US]; 995 Benecia Avenue, Sunnyvale, CA 94086 (US). (72) Inventors: RIEB, Dale; 1085 Tasman Drive #511, Sunnyvale, CA 94089 (US). FOSSE, Lorna; 96 Van Cott Court, San Jose, CA 95127 (US). POMERANZ, Mark, L.; 110 Ann Arbor Court, Los Gatos, CA 95032 (US). (74) Agents: STALLMAN, Michael, A. et al.; Limbach & Limbach L.L.P., 2001 Ferry Building, San Francisco, CA 94111 (US).			(81) Designated States: European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).  Published With international search report.

(54) Title: COMBINATION OF LINEAR ABLATION AND COOLED TIP RF CATHETERS



(57) Abstract

An apparatus for ablating body tissue is provided that is particularly adapted for creating both linear and point lesions in the endocardium. The apparatus comprises an elongate tubular member (26) having a tip electrode (32) and an ablation section (34) mounted thereon. The ablation section (34) includes one or more spaced electrodes, a fluid permeable foam material (46), and a fluid impermeable covering (48) having a plurality of holes (50) formed in it. The flow of conductive fluid to the ablation section during use allows contact to be maintained between the electrodes and the tissue to be ablated so as to minimize the formation of lesion breaks. The conductive fluid also serves to cool the tip electrode (32) during its use by delivering the fluid to the tip electrode before routing it to the ablation section. In preferred embodiments, use of a shape wire (66) and/or one or more pullwires (64) allows the inventive apparatus to be more easily manipulated during the ablation procedure.

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

## COMBINATION OF LINEAR ABLATION AND COOLED TIP RF CATHETERS

5

## BACKGROUND

The present invention generally relates to medical devices and methods for ablating living tissue. More particularly, the present invention relates to radio frequency ("RF") ablation catheters and methods for using the same to create lesions within the heart.

Atrial fibrillation is a condition in the heart in which abnormal electrical signals are generated in the endocardial tissue to cause irregular beatings of the heart. A proven protocol for successfully treating this condition is open heart surgery (sometimes referred to as the "maze" procedure) where several long (i.e. approximately 2-10 cm) lesions are created in the endocardium within the upper chambers of the heart ("atria"). These lesions block the flow of excess electrical impulses within the atria and allow the impulse from the sinus node to properly regulate heart contraction.

However, because open heart surgery is highly invasive and requires a lengthy patient recovery period, alternative methods for making lesions have been recently explored. One such alternative is the use of ablation catheters that includes one or more electrodes.

Typically, an ablation catheter is advanced into the heart via the patient's vessels. When the electrodes are placed in the desired position within the heart chamber, radio frequency ("RF") energy is supplied to the catheter thereby burning lesions into the endocardium.

Initial designs for ablation catheters generally comprised of an elongated shaft in which an electrode is mounted onto its distal end. Either point and linear lesions could be formed with these catheters by manipulating the placement of the tip. However, because of the tendency for the tip electrode to overheat and to lift off the tissue surface during ablation, creating suitable lesions using these catheters have been difficult.

New catheter designs attempted to mitigate these disadvantages. One improvement is the addition of a mechanism to cool the tip electrode during use to minimize the risk of embolism from overheated blood. Such cooled tip ablation catheters are described in U.S. Patent Nos. 5,423,811 and 5,545,161 both of which are incorporated in their entireties herein by reference. Although these catheters mitigate the overheating problem, the tendency for the tip electrode to form uneven linear lesions still remains.

Another improvement is the substitution of the tip electrode for a series of electrodes to form a linear ablation section on the catheter. Illustrative examples of such ablation catheters are described in pending U.S. Application No. 08/965,353 filed November 6, 1997, entitled "DEFLECTABLE LOOP DESIGN FOR A LINEAR LESION ABLATION APPARATUS" by inventors Mark L. Pomeranz, Troy J. Chapman, Scott Tedder,

Darren R. Sherman, and Steven C. Anderson, pending  
U.S. Application No. 08/680,426 filed on July 15,  
1996 entitled "SHAPABLE CATHETER USING EXCHANGEABLE  
CORE AND METHOD OF USE" by inventors Mark L. Pomeranz  
5 and Peter Park, and allowed U.S. Application No.  
08/611,656 filed on March 6, 1996 entitled "APPARATUS  
AND METHOD FOR LINEAR LESION ABLATION" by inventors  
Mark L. Pomeranz, Troy J. Chapman, Darren R. Sherman,  
and Mir Imran all of which are also incorporated in  
10 their entireties herein by reference.

These catheters facilitate the formation of suitable  
linear lesions by providing additional surface area  
for distributing RF energy. Although linear lesion  
catheters are a significant improvement over tip  
15 electrode catheters, breaks still occasionally occur  
due to the difficulty in maintaining sufficient  
contact between the ablation section and the tissue  
surface.

The impact of these breaks may be generally mitigated  
20 by forming point lesions at these sites in a follow  
up procedure. However, since linear ablation  
catheters are not generally suited to form point  
lesions, the follow up procedure typically requires  
removing the existing linear ablation catheter from  
25 the patient and substituting it for a tip electrode  
catheter. Because this catheter exchange may cause  
additional trauma to the patient as well as increase  
the overall length of the procedure, a need exists  
for an ablation catheter that combines the  
30 functionalities of both the tip electrode and the  
linear ablation section.

#### SUMMARY OF THE INVENTION

The present invention provides improved catheters for  
use in ablating tissue that allows for a greater

degree of control in the type and quality of the lesions that may be formed. The inventive catheters comprise an elongate tubular member having a tip electrode and an ablation section mounted thereon.

5 The ablation section includes one or more spaced electrodes, a fluid permeable foam material, and a fluid impermeable covering having a plurality of holes formed in it. Either point and linear lesions may be created by selectively and separately

10 controlling the delivery of current to either the tip electrode or the ablation section electrodes. The flow of conductive fluid to the ablation section during use allows contact to be maintained between the electrodes and the tissue to be ablated so as to

15 minimize the formation of breaks in the lesions. The conductive fluid also serves to cool the tip electrode during its use by delivering the fluid to the tip electrode before routing it to the ablation section. In preferred embodiments, use of a shape

20 wire and/or one or more pullwires provide additional maneuverability.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 is a side view of the preferred embodiment of an ablation catheter according to the present

25 invention.

Figure 2A is a more detailed view of the deflectable shaft region of the ablation catheter of Figure 1.

Figure 2B is a cross section of the main shaft at the designated plane B-B in Figure 2A.

30 Figure 2C is a cross section of the ablation section at the designated plane C-C in Figure 2A.

Figure 3A is an enlarged view of the ablation section without the foam and covering of the catheter of Figure 1.

5      Figure 3B-3E are cross sections at their respective designations of the ablation section in Figure 3A.

Figure 4 is an enlarged side view of the tip electrode of the subject ablation catheter.

Figure 5 is an alternative embodiment of the tip electrode.

10            **DETAILED DESCRIPTION OF THE DETAILED EMBODIMENTS**  
The present invention provides a combination ablation catheter that comprises a cooled tip electrode and a linear electrode region that allows an operator to have a greater degree of control in the type and  
15      quality of lesions that may be formed. The advantages of the combination ablation catheter of the present invention will be further described below.

Referring to Figure 1, inventive catheter 10  
20      generally comprises main shaft 20 having a distal end 22 and proximal end 24; deflectable shaft 26 also having a distal end 28 and proximal end 30; tip electrode 32; and ablation section 34 mounted on to deflectable shaft 26. As depicted by Figure 1,  
25      catheter 10 also may include handle 36 with deflection knob 38, fluid port 40, and female adapter 42 formed therein.

Proximal end 30 of deflectable shaft 26 and distal end 22 of main shaft 20 are contiguous so as to form  
30      one continuous shaft. The diameter and length of continuous shaft formed by deflectable shaft 26 and

main shaft 20 may be of any suitable size that may be required for an ablation procedure. In preferred embodiments, the length of deflectable shaft 26 is between about 3 cm to about 10 cm and the diameter of both deflectable shaft 26 and main shaft 20 is between about 2 mm and 5 mm.

Both main shaft 20 and deflectable shaft 26 may be made from the same material. Suitable materials include thermoplastic polymer, polyamide ether, polyurethane and other materials having similar properties that are known in the art.

Deflectable shaft 26 differs from main shaft 20 in that it is more flexible (i.e., less rigid or stiff). This difference in flexibility allows deflectable shaft 26 to deflect to any angle between 0 to about 180° from its non-deflected state, or assume various shapes without substantially affecting main shaft 20. As it will be explained in more detail below, the change in shape of deflectable shaft 26 may be a result of actuating a pullwire or the use of a removable core wire which is formed at least in part of memory wire. In preferred embodiments, the rigidity of main shaft 20 is between about 55 and about 80 durometers and the rigidity of deflectable shaft 26 is between about 30 and about 45 durometers.

Referring to Figures 2 and 3, ablation section 34 is mounted on at the distal half of deflectable shaft 28 and includes one or more spaced electrodes 44, fluid permeable deformable member or foam support segment 46, and a fluid impermeable covering 48 having a plurality of holes or openings 50 formed in it. In preferred embodiments, ablation section 34 has a length of approximately between 1 cm and 5 cm and an



outer diameter of approximately between 2 mm and 5 mm.

Although covering 48 may contain any number of holes in any pattern, arranging the holes along the side which will be positioned against the ablation site at a density of approximately between 20 holes/cm and 60 holes/cm is preferred. In especially preferred embodiments, covering 48 contains four rows of 0.007" diameter holes spaced 1.0 mm apart over the length of the active region of the ablation section.

Foam support segment 46 at least partially surrounds ablation section electrodes 44 and is itself enclosed within fluid impermeable covering 48 which is preferably formed of a polymeric material. Foam support segment 46 may be formed from any suitable material such as an open cell polyurethane, cotton-like material, open-cell sponge, hydrogels, or other fluid permeable compressible materials.

During use, RF energy is delivered to ablation section electrodes 44 while conductive fluid, such as saline, is simultaneously delivered to ablation section 34. The conductive fluid contacts electrodes 44 and flows out foam support segment 46 through holes 50 in covering 48. Means for delivering current and fluid to ablation section 34 will be described in detail further below.

Contact between ablation section 34 and the tissue to be ablated is enhanced in two ways. First, being deformable, foam support segment 46 can conform to the contours of the tissue to be ablated. The contact between ablation section 34 and the tissue to be ablated is further assisted by the positive pressure from the infusion of conductive fluid

through foam support segment 46. Second, because the fluid is conductive, the flow of fluid serves to couple the RF energy from ablation section electrodes 44 to the tissue even if direct contact between the electrodes and the tissue is not maintained. Moreover, the conductive liquid also serves to cool ablation section electrodes 44 thus making the procedure safer by decreasing the likelihood of thrombus formation from heated blood.

As shown in greater detail in Figure 4, tip electrode 32 is attached to the most distal end 28 of deflectable shaft 26 by conventional means known in the art and includes chamber 52 formed therein. In preferred embodiments, tip electrode 32 has a rounded distal end and is between about 2 mm and 5 mm in length. As it will be further described below, tip electrode 32 is also cooled by the flow of fluid. Although tip electrode 32 may optionally include holes to allow fluid to flow therethrough as with ablation section 34, preferred embodiments deliver fluid to tip electrode chamber 52 and then remove this fluid by diverting it back through the catheter shaft in the opposite direction.

Although alternate means may be used, the internal region of the rounded distal end of tip electrode 32 is filled with conductive material 54 such as solder which acts to distribute current from lead wire 56 (via exposed wire 57) to the tip of electrode 32. A device for monitoring the temperature of tip electrode 32 may optionally be included, such as thermocouple 58 depicted in Figure 4. However, it will be necessary to isolate thermocouple 58 from conductive material 54 by any known methods such as enclosing it in nonconductive pipe 60 filled with nonconductive adhesive 62.

Catheter maneuverability is provided by the use of one or more pullwires and/or one or more removable core wires, some of which are formed at least in part from memory or shape wire such as Nitinol. Referring back to Figure 1, pullwire 64 (not depicted in this illustration) is attached to proximal portion (at or near end 30) of deflectable shaft 26 by conventional means and the other end is attached to deflectable knob 38 in handle 36. Alternatively, pullwire 64 may be attached within chamber 52 of tip electrode 32.

Pullwire 64 is actuated by slidably retracting deflection knob 38 along handle 36 in the direction shown by arrow A in Figure 1. The retraction of knob 38 causes deflectable shaft 26 to be deflected. When knob 38 is moved back along handle 36 in a direction opposite arrow A, deflectable shaft 26 returns to its nondeflected state. Alternate pullwire actuation means may also be used. Although this embodiment uses only one pullwire, additional deflection points may be created by using a plurality of pullwires.

Exchangeable core wires, some of which are made at least in part of shape or memory wire, may be used either in conjunction with, or as an alternative to the one or more pullwires. Typically, only the portion of the core wire which will be carried within deflectable shaft 26 (so as to assume the desired shape) needs to be formed from memory or shape wire. As suggested by Figure 1, core wire 66 is slidably disposed within a lumen that extends from distal end 28 of deflectable shaft 26 through the proximal portion of main shaft 20, and exits out of main shaft 20 at an opening therein. Core wire 66 preferably has a substantially flat cross-section (i.e. rectangular or oblong) to prevent the core wire from rotating about its longitudinal axis during use. In

addition, means for stiffening the portion of core wire 66 which diverges from main shaft 20 is also preferred, like attaching core wire 66 to a more rigid wire 68. This facilitates the insertion and removal of core wire 66 from the catheter shaft.

During use, either a catheter without a core wire or a substantially straight core wire is used to advance the instrument through a patient's vessels. However, because of increased flexibility for threading through a patient's vessels, use of a catheter without a core wire inserted therein is generally preferred. When it is desired for deflectable shaft 26 to assume a certain shape, then an appropriate core wire that will form this shape is inserted into the catheter shaft via the opening in the proximal end 24 of main shaft 20. Removing existing core wire 66 and replacing it with one assuming a different shape may occur several times during the ablation procedure.

If both pullwire(s) and exchange core wire are to be used in the same catheter, then it is preferred that the distal ends of the pullwires all terminate in the shaft proximal to the section in deflectable shaft 26 that will assume the desired core wire shape.

Any suitable means may be used to deliver current to ablation section electrodes 44 and tip electrode 32. In preferred embodiments, current may be delivered to tip electrode 32 without also delivering current to ablation section electrodes 44 and vice versa. This is typically achieved by using a switch box linked to catheter 10 via female port 42 that allows RF energy to be directed from an RF source to either the tip electrode lead wire(s) or the ablation electrode lead wires or if desired, both. The RF source may be any

conventional RF generator such as Model 8002 or 8004 RF Generator by Cardiac Pathways, Inc. (Sunnyvale, California).

5 Similarly, any conventional means may be used to deliver conductive fluid to ablation section 34, and to deliver and remove cooling fluid to and from tip electrode 32. For the sake of efficiency, it is preferred that the conductive fluid is also used as the cooling fluid for tip electrode 32.

10 Although any number of lumens may be used to accomplish delivering current and fluid to the appropriate sites, minimizing the number of lumens necessary for the task is generally preferred for manufacturing expediency. In preferred embodiments,  
15 main shaft 20 and deflectable shaft 26 include four lumens therein. These lumens are illustrated in Figures 2B-2C and Figures 3B-3E.

Figure 2B illustrates a cross section of main shaft 20. The four lumens include lumen 70 for core wire  
20 66; lumen 72 for tip electrode lead 56 and thermocouple leads 59; lumen 74 for ablation electrode leads 51; and lumen 76 for pullwire 64. As it will be further explained below, lumen 72 is also  
25 used to deliver fluid from fluid port 40 to chamber 52 of tip electrode 32 and the distal portion of lumen 74 is used to route fluid from chamber 52 to ablation section 34.

A stainless steel braid 78 is preferably embedded in the wall of main shaft 20 by conventional means known  
30 in the art. The inclusion of the braid improves the torque characteristics of main shaft 20 and thus makes the shaft easier to maneuver through a patient's vessels and heart. To minimize the

possibility for tissue trauma as the ablation catheter is fed through a patient's blood vessels and heart during use, wire braid 78 is preferably absent from deflectable shaft 26. This leaves the catheter tip sufficiently flexible to yield when advanced against obstacles within the vessels and heart.

Figure 2C is a cross section of ablation section 34. Lumens 70, 72, and 74, and their respective contents are as previously described. Lumen 76 exists but is empty because pullwire 64 terminated at some point within the proximal portion of deflectable shaft 26. However, as previously described, if the pullwire terminated within chamber 52 of tip electrode 32 instead of the proximal portion of deflectable shaft 26 then pullwire 64 would be present in lumen 76. Additionally, Figure 2C shows foam support segment 46 and fluid impermeable covering 48.

Figures 3B-3E are additional cross sections of ablation section 34 which serve to illustrate the preferred coupling between the means for cooling tip electrode 32 and the means for delivering conductive fluid to ablation section 34. As described previously, lumen 72 delivers fluid from fluid port 40 to chamber 52 of tip electrode 32 (as well as carry tip electrode lead 56 and thermocouple lead 59).

Figure 3E is a cross section which is closest to tip electrode 32. Lumen 70 carrying core wire 66 is not seen since it terminated at a position proximal to this cross section. Lumen 74 which carries ablation section electrode leads 51 is empty since the leads terminated previously at their respective electrodes.

Conductive cooling fluid is delivered to chamber 52 via lumen 72. When the fluid reaches chamber 52, fluid fills the chamber and is routed back away from tip electrode 32 through lumen 74. When an  
5 alternative embodiment for the tip electrode is used, such as that depicted by Figure 5, then cooling fluid will also exit out from tip electrode 32 from holes 80. In either embodiments, fluid is not able to flow back through lumen 72 because of the positive  
10 pressure created by the inflowing fluid. This continuous flow of fluid results in cooling tip electrode 32 during use.

From tip electrode chamber 52, conductive fluid is routed back through the catheter shaft towards  
15 ablation section 34 via lumen 74 (which also carries ablation electrode leads 51). Openings (not depicted) in lumen 74 within ablation section 34 allows fluid to flow from lumen 74 through foam support member 46 and out holes 50 in covering 48.  
20 The flow of fluid does not continue through lumen 74 toward main shaft 20 because as depicted by Figure 3B, lumen 74 is closed off at a point proximal to ablation section 34.

It should be understood that various features of the preferred embodiments may be used in any combination  
25 with one another. For example, catheter 10 may include only pullwire 64 and not include exchangeable core wire 66. Alternatively, the means for delivering conductive fluid to ablation section 34  
30 may be independent from the means for delivering cooling fluid to and for removing cooling fluid from tip electrode 32.

Moreover, although the present invention has been described with reference to preferred embodiments, it

should be appreciated that these embodiments are for purposes of illustration only and are not intended to limit the scope of the appended claims.



What is claimed is:

1. An apparatus for ablating body tissue,  
comprising:
  - an elongate tubular member having a distal end;
  - 5 a tip electrode attached to the distal end of  
the elongated member;
  - an ablation section having at one or more  
electrodes carried on the elongate tubular member;
  - a fluid permeable deformable member at least  
10 partially covering the ablation section electrodes;  
means for delivering current to the tip  
electrode;
  - means for delivering current to the ablation  
section electrodes; and,
  - 15 means for delivering conductive fluid through  
the deformable member in the ablation section such  
that when the electrodes are placed adjacent to the  
body tissue, the fluid creates a conductive path  
between the ablation section electrodes and the  
20 tissue.
2. The apparatus of claim 1 further comprising a  
core wire carried within the elongate tubular member.
3. The apparatus of claim 2 wherein the core wire  
is formed at least in part of memory wire and the  
25 elongate tubular member includes an opening at its  
proximal end, such that the core wire is slidably  
disposed in and removable from the elongate tubular  
member through the proximal end opening of the  
elongate tubular member.
- 30 4. The apparatus of claim 1 wherein the means for  
delivering the current to the tip electrode is  
independent from the means for delivering the current  
to the ablation section such that current may be

separately delivered to either the tip electrode or the ablation section electrodes.

5. The apparatus of claim 1 wherein the tip electrode includes a chamber formed therein and the apparatus further comprises:

means for delivering cooling fluid to the tip electrode chamber wherein the cooling fluid cools the tip electrode during use and

means for withdrawing the cooling fluid from the tip electrode chamber after its use.

6. The apparatus of claim 5 wherein the cooling fluid is also conductive and the means for delivering conductive fluid to ablation section is coupled to the means for removing cooling fluid from the tip electrode chamber.

7. The apparatus of claim 6 wherein the cooling fluid is withdrawn from the tip electrode chamber by delivering the cooling liquid through the deformable member in the ablation section.

8. The apparatus of claim 5 wherein the tip electrode chamber contains one or more openings for allowing at least some portion of the cooling liquid to exit out the tip electrode chamber while the tissue is being ablated.

9. The apparatus of claim 1 wherein the deformable member includes a layer of foam over the ablation section electrodes.

10. The apparatus of claim 9 further comprising a covering on the deformable member, the covering formed of a material substantially impermeable to

fluid, the covering including at least one opening sized to allow passage of fluid out of the covering.

11. The apparatus of claim 1 wherein the tip electrode includes a means for measuring temperature.

5 12. The apparatus of claim 1 wherein the distal end of the elongate tubular member is bendable and wherein the apparatus further comprises:

10 a pullwire extending through the elongate tubular member, the pullwire having a fixed first end and a moveable second end for adjusting the bend of the distal end of the elongate tubular member.

13. The apparatus of claim 1 further comprising a plurality of pullwires and a plurality of means for actuating the pullwires.

15 14. A catheter for ablating body tissue, comprising:  
an shaft having a bendable distal section and a distal end;

a tip electrode having a chamber formed therein and attached to the distal end of the shaft;

20 an ablation element carried on the distal section of the shaft; and,

means for delivering current to the tip electrode;

25 means for delivering current to the ablation element;

means for delivering fluid to the tip electrode chamber to cool the tip electrode during use; and,

means for withdrawing the fluid from the tip electrode.

30 15. The catheter of claim 14 wherein the tip electrode include a means for measuring temperature.

16. The catheter of claim 14 wherein the means for delivering the current to the tip electrode is independent from the means for delivering the current to the ablation section such that current may be  
5 separately delivered to either the tip electrode or the ablation section electrodes.

17. The catheter of claim 14 further comprising:  
a pullwire extending through the shaft member,  
and having a fixed distal end and a moveable proximal  
10 end for adjusting the bend of the distal end of the shaft member.

18. The catheter of claim 14 further comprising:  
a fluid permeable deformable member at least  
partially covering the ablation element and  
15 means for delivering conductive fluid through the deformable member.

19. The catheter of claim 14 wherein the cooling fluid is conductive.

20. The catheter of claim 19 further comprising:  
20 a fluid permeable deformable member at least partially covering the ablation element and wherein the means for withdrawing the fluid from the tip electrode is by delivering the fluid through the deformable member.

21. The catheter of claim 18 further comprising a  
25 covering on the deformable member, the covering formed of a material substantially impermeable to fluid, the covering including at least one opening sized to allow passage of fluid out of the covering.

22. The catheter of claim 14 further comprising a  
30 core wire carried within the shaft member.

23. The catheter of claim 22 wherein the core wire is formed at least in part of memory wire and the shaft member includes an opening at its proximal end, such that the core wire is slidably disposed in and removable from the shaft member through the proximal end opening of the shaft.

24. The catheter of claim 14 wherein the tip electrode chamber includes at least one opening sized to allow fluid to exit from the tip electrode during ablation.

25. A multipurpose device for ablating tissue comprising:

an elongated catheter having a fluid carrying lumen extending from the proximal to the distal end thereof;

a first electrode located at the distal end of the catheter, the first electrode being thermodynamically coupled to fluid carried in the lumen;

a plurality of second electrodes located near but spaced apart from the distal end of the catheter, the second electrodes being covered by elongated fluid permeable material and coupled to the lumen so that fluid can pass out of the lumen, through the fluid permeable material and to the tissue; and

means for delivering current to the first and second electrodes.

26. A method for ablating tissue using a catheter having a tip electrode located on its distal end and a plurality of second electrodes located near but spaced apart from the tip electrode, comprising:  
forming at least one linear lesion using the plurality of second electrodes and then

forming at least one point lesion using the tip electrode.

5 27. The method as in claim 26 wherein the point lesion is formed in the same site as the linear lesion.

28. A method for ablating tissue using a catheter having a tip electrode located on its distal end and a plurality of second electrodes located near but spaced apart from the tip electrode, comprising:

10 forming at least one point lesion using the tip electrode and then

forming at least one linear lesion using the plurality of second electrodes.



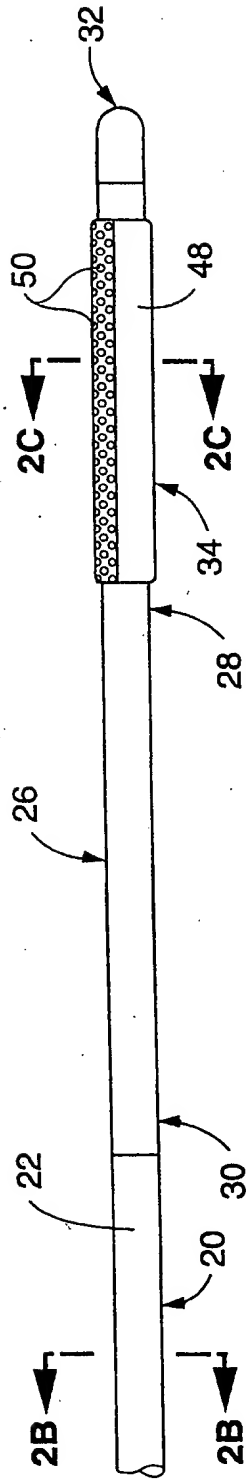


FIG. 2A

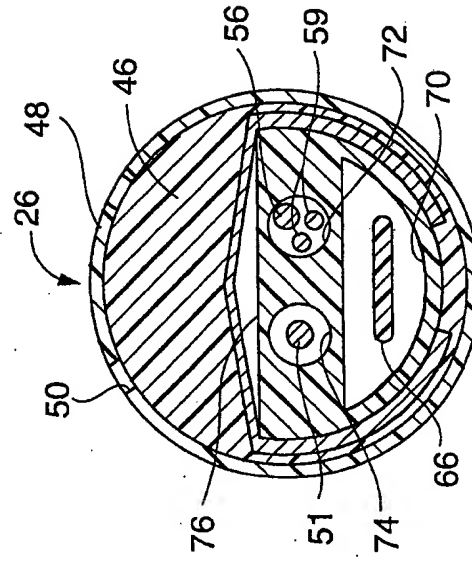


FIG. 2C

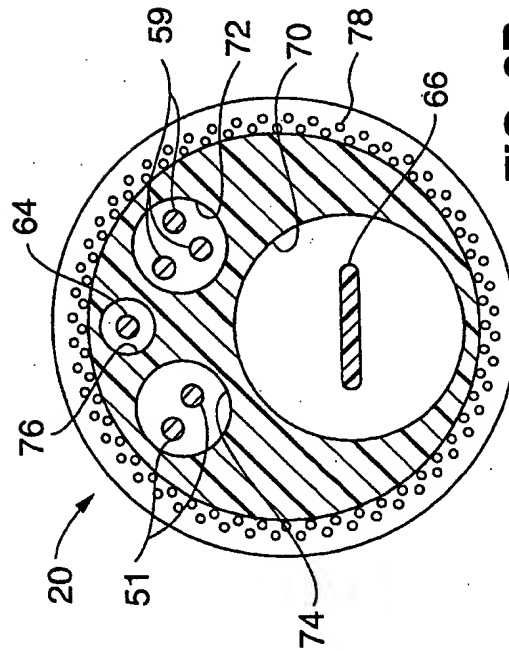


FIG. 2B



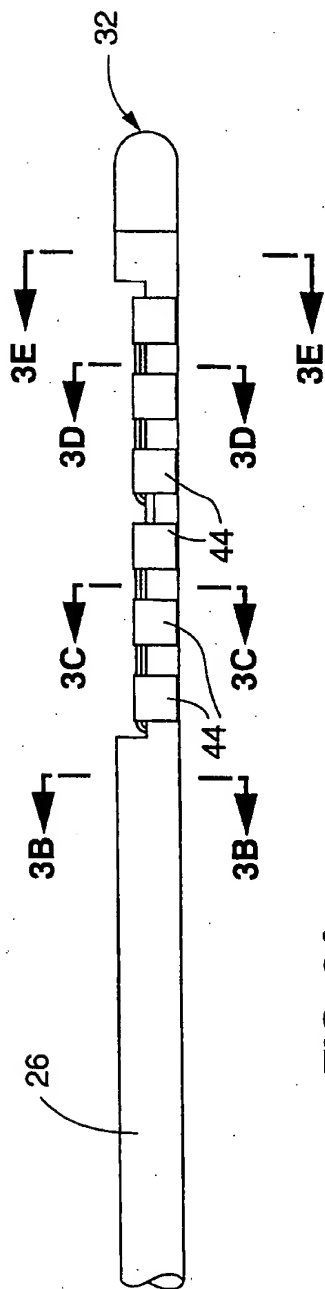


FIG. 3A

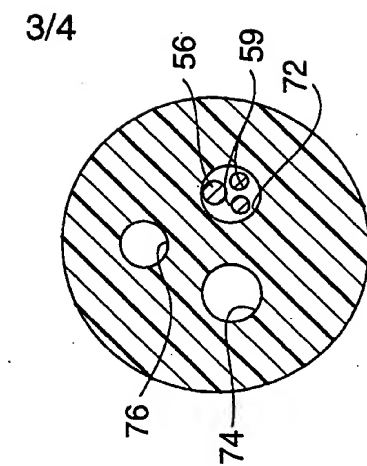


FIG. 3B

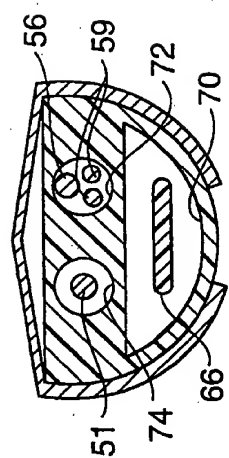


FIG. 3C

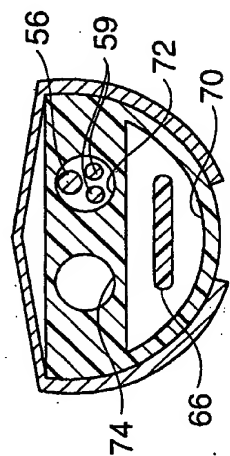


FIG. 3D

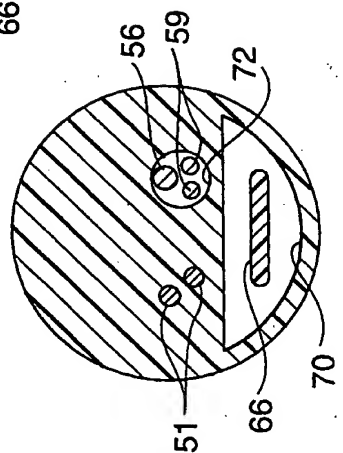
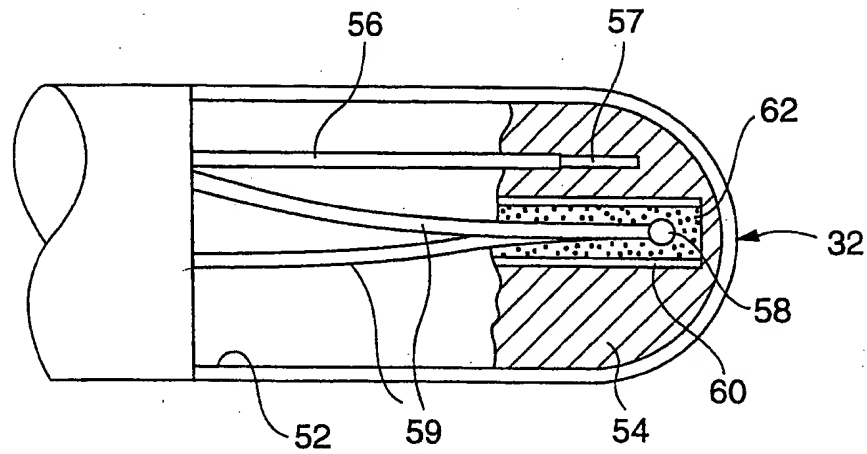
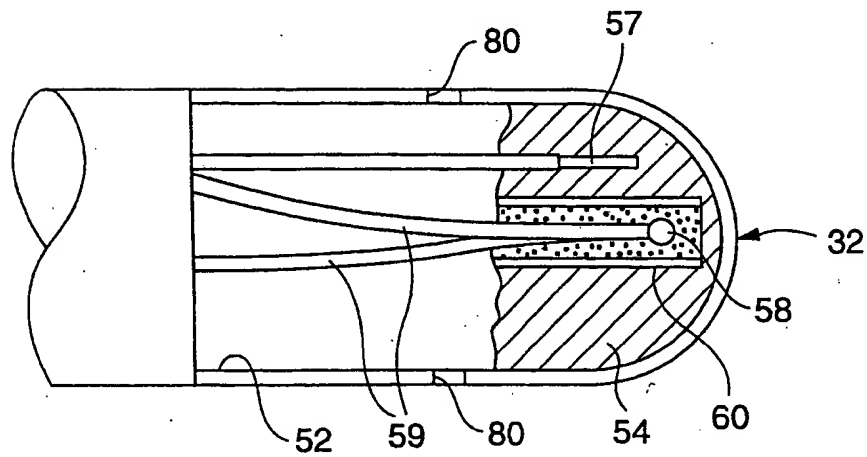


FIG. 3E

4/4



**FIG. 4**



**FIG. 5**

# INTERNATIONAL SEARCH REPORT

Internal Application No  
PCT/US 99/00567

A. CLASSIFICATION OF SUBJECT MATTER		
IPC 6	A61B5/042	A61N1/05 G01B7/28 A61M25/00 A61M25/01
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED		
Minimum documentation searched (classification system followed by classification symbols)		
IPC 6 A61B A61N G01B A61M		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 423 811 A (IMRAN MIR A ET AL) 13 June 1995 see column 2, line 38-57	14,15, 19,24 1,2,5,6, 8-11,17, 18, 20-22,25
Y	see column 3, line 8 - column 4, line 46 see column 5, line 45-57 see column 6, line 14 - column 7, line 4 see figures 1,2,4	
Y	WO 98 02201 A (CARDIAC PATHWAYS CORP) 22 January 1998 see page 4 see page 6, line 24 - page 7, line 22 see page 11, line 19 - page 13, line 26 see figures 1,2,9-12	1,2,5,6, 8-11,18, 20-22,25
-/-		
<input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex.		
* Special categories of cited documents : "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "Z" document member of the same patent family		
Date of the actual completion of the international search		Date of mailing of the international search report
28 April 1999		11.05.99
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer  Bichlmayer, K-P

# INTERNATIONAL SEARCH REPORT

Internal Application No  
PCT/US 99/00567

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 5 676 693 A (LAFONTAINE DANIEL MARC) 14 October 1997 see column 2, line 54 - column 3, line 2 see figure 3 ---	1,25
Y	WO 97 32525 A (CARDIAC PATHWAYS CORP) 12 September 1997 see page 3, line 19 - page 4, line 34; figures 12A-E ---	18,20,21
A	EP 0 395 098 A (TOKIN CORP ;TERUMO CORP (JP)) 31 October 1990 see abstract ---	3,23
A	US 5 730 127 A (AVITALL BOAZ) 24 March 1998 see column 6, line 50-57 ---	4,16
A	EP 0 689 851 A (CORDIS EUROP) 3 January 1996 see the whole document ---	12 17
A	US 4 570 354 A (HINDES CHARLES S) 18 February 1986 see column 2, line 45-60 -----	13 17

1.

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US 99/00567

### Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 26-28  
because they relate to subject matter not required to be searched by this Authority, namely:  
Rule 39.1(iv) PCT - Method for treatment of the human or animal body by surgery
2. ☐ Claims Nos.:  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

### Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.  
☐ No protest accompanied the payment of additional search fees.

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/00567

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5423811 A	13-06-1995	US 5348554 A	20-09-1994
		US 5658278 A	19-08-1997
		AU 673015 B	24-10-1996
		AU 5201993 A	16-06-1994
		CA 2110309 A	02-06-1994
		EP 0608609 A	03-08-1994
		JP 2562861 B	11-12-1996
		JP 6254103 A	13-09-1994
		US 5697927 A	16-12-1997
		US 5545161 A	13-08-1996
WO 9802201 A	22-01-1998	US 5882346 A	16-03-1999
		AU 3153997 A	09-02-1998
US 5676693 A	14-10-1997	EP 0765178 A	02-04-1997
		JP 10501439 T	10-02-1998
		WO 9534346 A	21-12-1995
		DE 4338758 A	19-05-1994
		US 5584872 A	17-12-1996
WO 9732525 A	12-09-1997	US 5800482 A	01-09-1998
		EP 0884978 A	23-12-1998
EP 0395098 A	31-10-1990	JP 1716271 C	27-11-1992
		JP 2289265 A	29-11-1990
		JP 4002273 B	17-01-1992
		JP 1716272 C	27-11-1992
		JP 2289266 A	29-11-1990
		JP 4002274 B	17-01-1992
		JP 1736744 C	26-02-1993
		JP 2289267 A	29-11-1990
		JP 4008065 B	13-02-1992
		AU 623006 B	30-04-1992
		AU 5451590 A	01-11-1990
		DE 69007841 D	11-05-1994
		DE 69007841 T	11-08-1994
		KR 9405307 B	16-06-1994
		US 5069226 A	03-12-1991
US 5730127 A	24-03-1998	US 5687723 A	18-11-1997
		US 5487385 A	30-01-1996
		US 5842984 A	01-12-1998
		AU 685559 B	22-01-1998
		AU 1265495 A	19-06-1995
		CA 2177982 A	08-06-1995
		EP 0731665 A	18-09-1996
		JP 9506017 T	17-06-1997
		WO 9515115 A	08-06-1997
EP 0689851 A	03-01-1996	NL 9401107 A	01-02-1996
		US 5674197 A	07-10-1997
US 4570354 A	18-02-1986	NONE	